### THE NT OF DEPT.

### DEPARTMENT OF THE AIR FORCE

### 59TH MEDICAL WING (AETC) LACKLAND AIR FORCE BASE TEXAS

18 MAR 2016

MEMORANDUM FOR SGVT

ATTN: LT COL MICHAEL R DAVIS

FROM: 59 MDW/SGVU

SUBJECT: Professional Presentation Approval

- 1. Your paper, entitled <u>Locally Applied Enzyme Activated Tacrolimus Eluting Hydrogels Significantly Delay the Onset of Acute Rejection of VCA Grafts presented at 2016 Society of Military Surgeons, Boston, MA, 17-19 March 2016</u> with MDWI 41-108, and has been assigned local file #16139.
- 2. Pertinent biographic information (name of author(s), title, etc.) has been entered into our computer file. Please advise us (by phone or mail) that your presentation was given. At that time, we will need the date (month, day and year) along with the location of your presentation. It is important to update this information so that we can provide quality support for you, your department, and the Medical Center commander. This information is used to document the scholarly activities of our professional staff and students, which is an essential component of Wilford Hall Ambulatory Surgical Center (WHASC) internship and residency programs.
- 3. Please know that if you are a Graduate Health Sciences Education student and your department has told you they cannot fund your publication, the 59th Clinical Research Division may pay for your basic journal publishing charges (to include costs for tables and black and white photos). We cannot pay for reprints. If you are 59 MDW staff member, we can forward your request for funds to the designated wing POC.
- 4. Congratulations, and thank you for your efforts and time. Your contributions are vital to the medical mission. We look forward to assisting you in your future publication/presentation efforts.

LINDA STEEL-GOODWIN, Col, USAF, BSC Director, Clinical Investigations & Research Support

hinder Steel-Goodwin

PROCESSING OF PROFES	SIONAL MEDICAL R	RESEARCH/TECHNICAL	PUBLICATION	IS/PRESENTATIONS		
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i. PROTOCOL TITLE: (NOTE: For each new release of medical research or lechnical information as a publication/presentation, a new 59 MDW Form 3039 must be submitted for review and approval.)						
Vascularized Composite Allotransplantation (VCA) in Swine (Sus scrofa) for Optimization of Reconstruction of Battlefield Injuries Using the						
6. TITLE OF MATERIAL TO BE PUBLISHED OR PRESENTED:						
Locally Applied enzyme activated tacrolimus eluting hydrogels significantly delay the onset of acute rejection of VCA grafts						
7. FUNDING RECEIVED FOR THIS STUDY?	YES NO FUN	DING SOURCE 59MDW ST				
8. DO YOU NEED FUNDING SUPPORT FOR PUBLICATION PURPOSES: YES NO						
9. IS THIS MATERIAL CLASSIFIED? TYES NO						
10. IS THIS MATERIAL SUBJECT TO ANY L AND DEVELOPMENT AGREEMENT (CRAD/ YES X NO NOTE: If the arower b YE	), MATERIAL TRANSFER	RAGREEMENT (MTA), INTELL	ECTUAL PROPERT	TY RIGHTS AGREEMENT ETC		
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11b. PUBLISHED ABSTRACT (List intended journal.)						
11c. POSTER (To be demonstrated at meeting: name of meeting, city, state, and date of meeting.)						
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12. EXPECTED DATE WHEN YOU WILL NE NOTE: All publications/presentations and				DITC OT NO		
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March 18, 2016						
13. 59 MDW PRIMARY POINT OF CONTACT (Last Namo, First Namo, M.I., email) 14. [			14. DUTY PHONE/PAGER NU	MBCR		
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15. AUTHORSHIP AND CO-AUTHOR(S) LI			EUCE EVIDOI	INSTITUTION (II and 50 b)	DIAN	
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b. Fries Charles A	0-4	59MDW ST				
c. Wu Kevin	VOL	59MDW ST				
d. Wang Lin C	0-3	59MDW ST				
e. Gorantla Vijay S	VOI.	59MDW ST				
f. Davis Michael R	0-5	59MDW ST				
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Christopher Carwile, TSgt/E-6, NCOIC, PA	STEWART.1280477229  STEWART.400477229  STEWART.400477229	18 March 2016			
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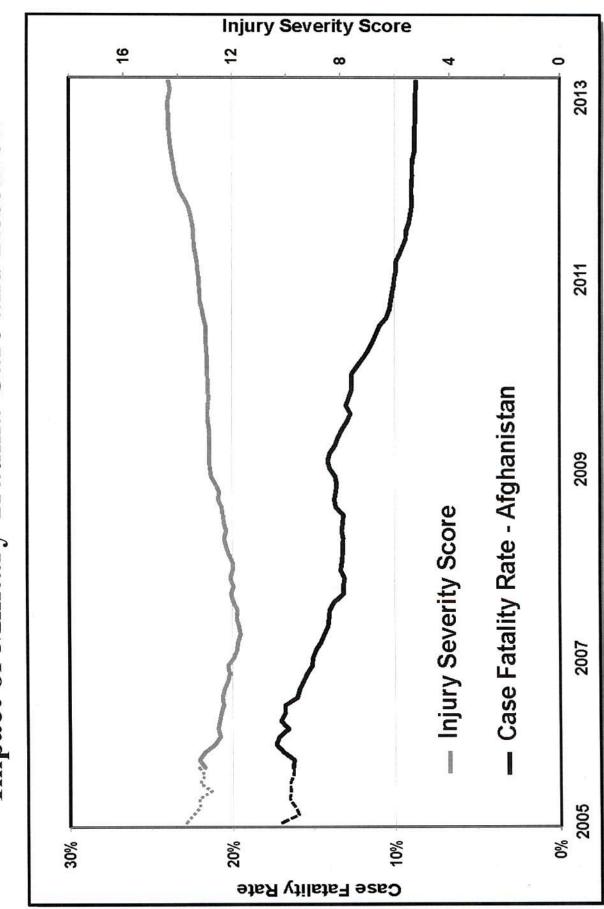
\* This person should be the frimary Author on Form 3039 General Surgery Resident, US Army Institute of Surgical Research, RESTOR Program, 59th Medical Wing ST CPT Renford Cindass Jr., MD

Deputy Commander, USA Institute of Surgical Research 59MDW Science and Technology Office Lt Col Michael R. Davis, MD, FACS San Antonio, TX, USA RESTOR Program

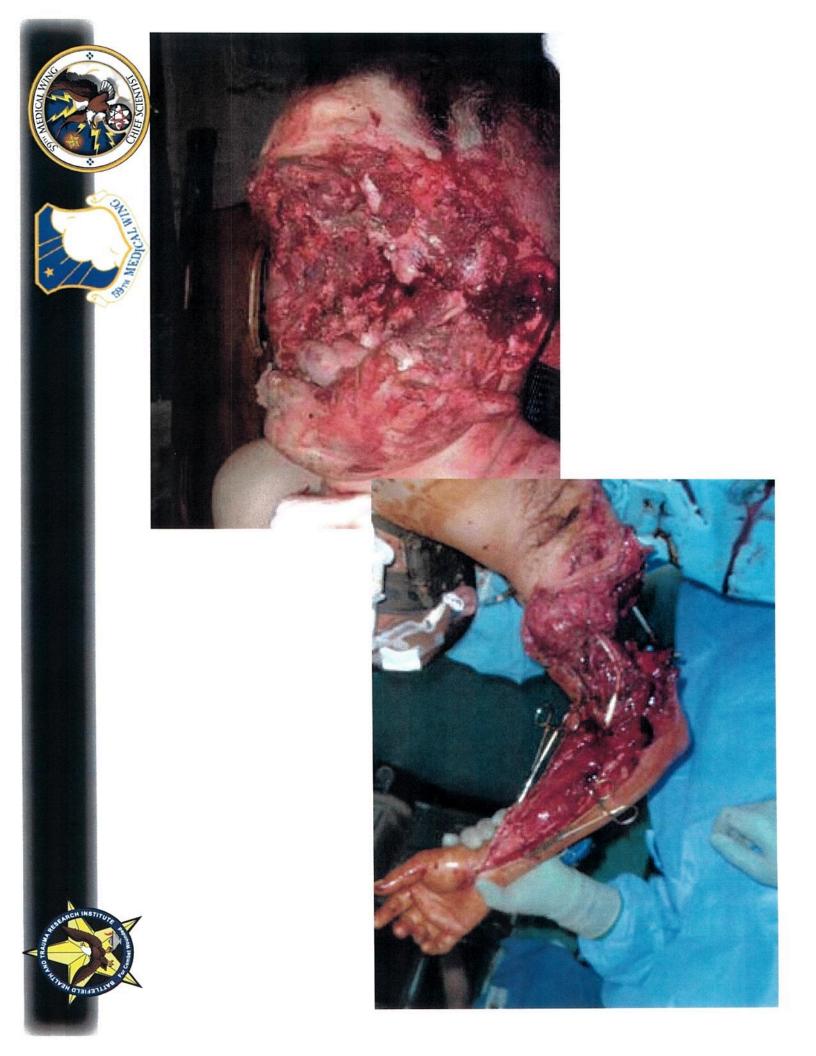




The opinions or assertions contained herein are the private views of the author and are not to be construed as official or as reflecting the views of the Department of Defense. The experiments reported herein were conducted according to the principles set forth in the National Institute of Health Publication No. 80-23, Guide for the Care and Use of Laboratory Animals and the Animal Welfare Act of 1966, as amended









New rung on the reconstructive ladder

Vascularized composite allotransplantation

Free tissue transfer eg, latissimus dorsi flap

Regional flaps
eg, posterior interosseous

Local flap eg, rotational/transposition

Skin graft

Secondary closure

Primary closure





Multiple types of tissues are transplanted as a single functional

Replaces like with like and restores form and function

Eliminates autologous donor site morbidity and minimizes the need for multiple reconstructive procedures



- Current challenges and limitations
- A life-enhancing but not a life-saving procedure
- Requires lifelong systemic immunosuppression
- Opportunistic infections: 88%
- Metabolic complications: 70%
- $-\ge 1$  episode of acute rejection within 1st year: 85%
- Limited to highly motivated patients
- Limited donor pool



## Background

A novel model of VCA

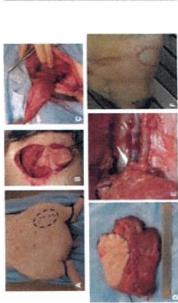
Evaluation of a drug eluting hydrogel



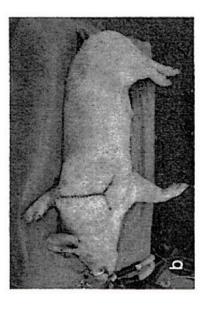


## A New Model

- Small animal models are technically challenging and lack immunologic maturity
- Previous orthotopic models in non-human primates and canines are no longer in use
- Currently only heterotopic swine models exist







Hettiaratchy, S., Melendy, E., Randolph, M. A., Coburn, R. C., Neville, D. M., Sachs, D. H., et al. (2004). Tolerance to composite tissue allografts across a major histocompatibility barrier in miniature swine.

Leto Barone, A. A., Leonard, D. A., Torabi, R., Mallard, C., Glor, T., Scalea, J. R., et al. (2013). The gracilis myocutaneous free flap in swine: an advantageous preclinical model for vascularized composite allograft transplantation research. Microsurgery, 33(1), 51–55

Kiermeir, D. M., Meoli, M., Müller, S., Abderhalden, S., Vögelin, E., & Constantinescu, M. A. (2013). Evaluation of a porcine whole-limb heterotopic autotransplantation model. Microsurgery, 33(2), 141–147





## A New Model

Ethically acceptable

Reproducible

Genetically controlled animals

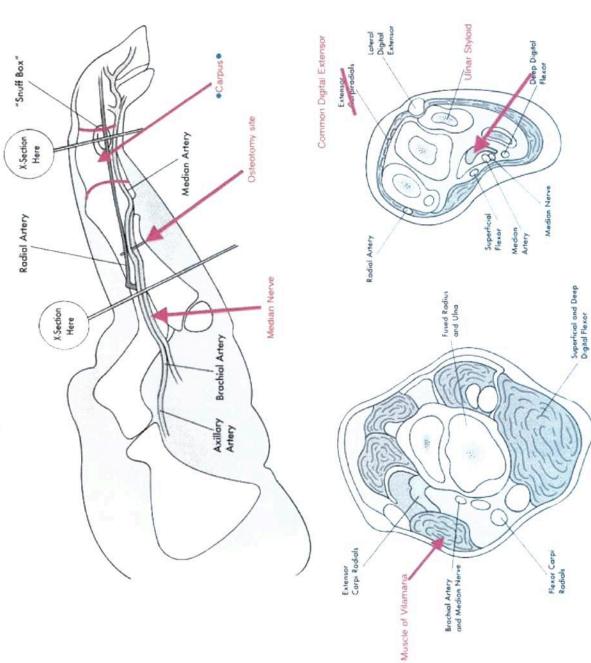
Orthotopic model to assess functionality

Evaluation of bone, tendon and nerve healing





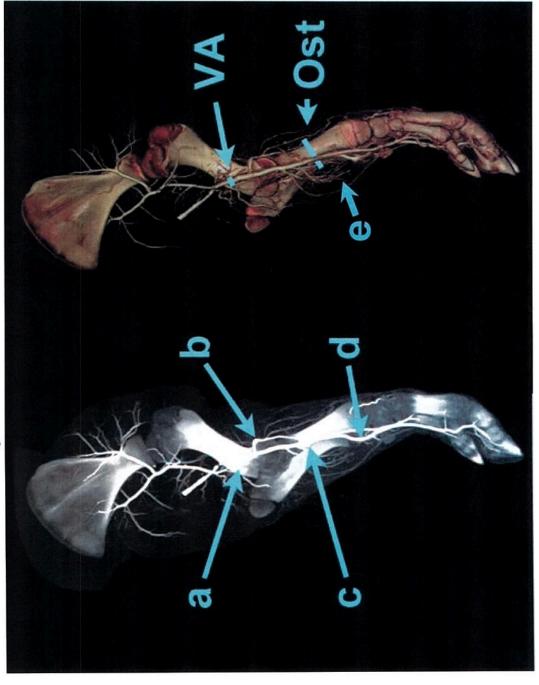
### Anatomy







### Anatomy



VA – Vascular Anastomosis Ost – Osteotomy Site

A – Axillary Artery B – Radial Artery

C – Interosseous Branch (of Fries)

D – Median Artery E – The Nest (of Lawson)



## 2 Weeks Post-op







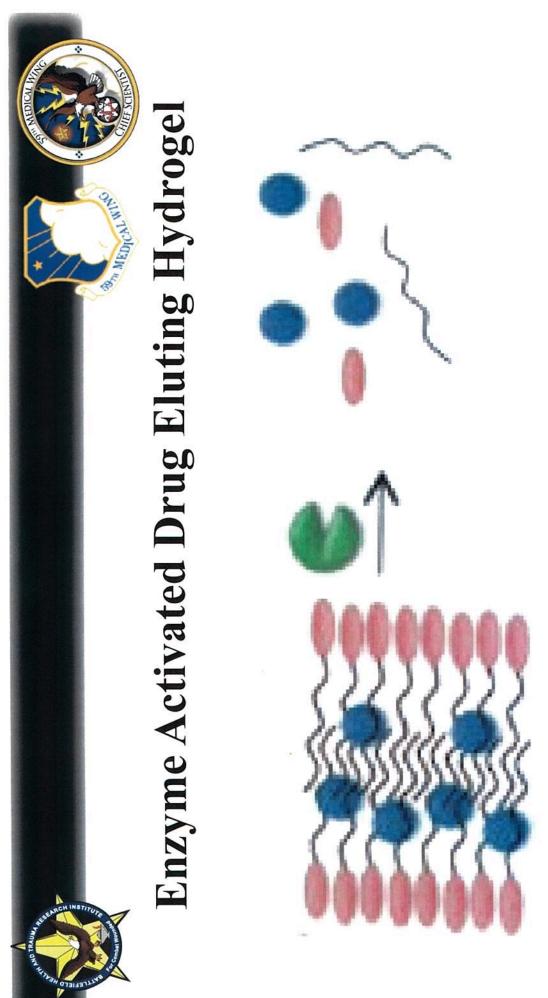
RESEARCH ARTICLE

TRANSPLANTATION

## A single localized dose of enzyme-responsive hydrogel improves long-term survival of a vascularized composite allograft

Thusitha Gajanayake, <sup>1,2</sup>\* Radu Olariu, <sup>1,2</sup>\* Franck M. Leclère, <sup>1,2</sup> Ashish Dhayani, <sup>3</sup> Zijiang Yang, <sup>4</sup> Anjan K. Bongoni, <sup>2,5</sup> Yara Banz, <sup>6</sup> Mihai A. Constantinescu, <sup>1,2</sup> Jeffrey M. Karp, <sup>4†</sup> Praveen Kumar Vemula, <sup>3†</sup> Robert Rieben, <sup>2†</sup> Esther Vögelin <sup>1,2</sup>

has considerable side effects and reduces the quality of life of VCA recipients. We loaded the immunosuppressive Currently, systemic immunosuppression is used in vascularized composite allotransplantation (VCA). This treatment drug tacrolimus into a self-assembled hydrogel, which releases the drug in response to proteolytic enzymes that are longed graft survival in a Brown Norway-to-Lewis rat hindlimb transplantation model, leading to a median graft survival of >100 days compared to 33.5 days in tacrolimus only-treated recipients. Control groups with no treatbodies and complement C3, revealed significantly reduced immune responses in the tacrolimus-hydrogel group compared with tacrolimus only. In conclusion, a single-dose local injection of an enzyme-responsive tacrolimushydrogel is capable of preventing VCA rejection for >100 days in a rat model and may offer a new approach for overexpressed during inflammation. A one-time local injection of the tacrolimus-laden hydrogel significantly proment or hydrogel only showed a graft survival of 11 days. Histopathological evaluation, including anti-graft anti-







### Methods



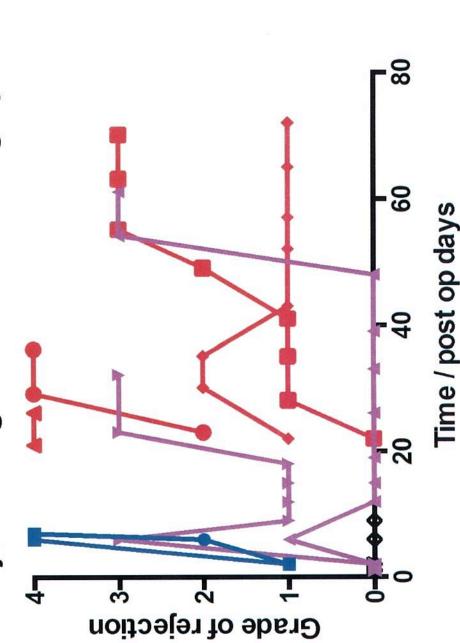
- Three groups
- Group 1: Controls no immunosuppression
- Group 2: High dose tacrolimus eluting hydrogel (84mg)
- Group 3: Low dose tacrolimus eluting hydrogel (49mg)
- 1 swine leukocyte antigen (SLA) donor-recipient mismatch
- No systemic immunosuppression
- Hydrogel injected in the subcutaneous layer following revascularization
- AST, LDH, CK, TNF-a, IL-6, myoglobin, and biopsies were assessed for signs of systemic toxicity and/or acute rejection
- End-point Banff grade 4 acute rejection or 100 days







# Rejection against time of allographs



High dose Tac

Control

Control

High dose Tac

Low dose Tac

Low dose Tac

Low dose Tac

Low dose Tac

POD2 Death

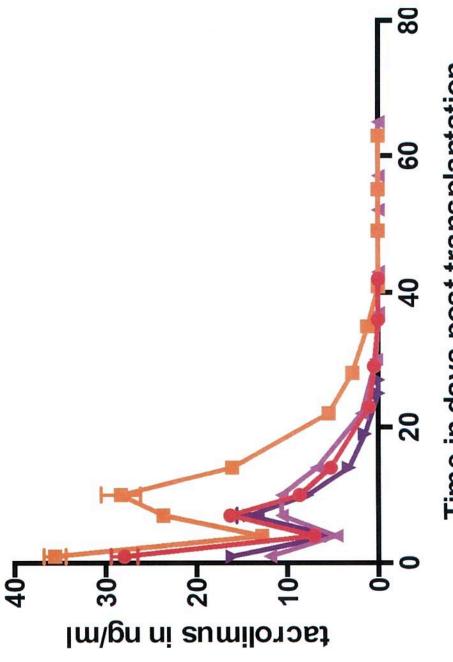
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### Results

# Tacrolimus systemic levels









## By MEDICAL WITH

## Conclusions

- The orthotopic model of swine VCA is an optimal model for investigating novel immunologic strategies
- Hydrogels are able to delay the onset of acute rejection with no gross safety concerns and without clinically detectable systemic levels of tacrolimus





## Future direction

Further hydrogel protocols to establish optimal dosing regimen and potential protocols for re-loading hydrogels

Increased survival duration to evaluate longer term rejection and side effects profile





## Thank you

### USAISR/59MDW

Lt Col Michael Davis Lt Col Dmitry Tuder

Dr Shari Lawson

CPT Lin Wang Dr Kevin Wu

Mr Raul Corpus

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